

Patent Claims

1. Process for the preparation of mono(fluoroalkyl)- or bis(fluoroalkyl)phosphoric acid, mono(fluoroalkyl) or bis(fluoroalkyl) phosphates and the corresponding phosphoranes thereof, comprising at least the reaction of a bis(fluoroalkyl)phosphinic acid or a (fluoroalkyl)phosphonic acid or a corresponding derivative or salt of these acids with anhydrous hydrogen fluoride.
- 10 2. Process according to Claim 1,
characterised in that use is made of a bis(fluoroalkyl)phosphinic acid or a corresponding derivative in which the two fluoroalkyl groups are identical or different.
- 15 3. Process according to Claim 1 or 2,
characterised in that use is made of a bis(perfluoroalkyl)phosphinic acid or a (perfluoroalkyl)phosphonic acid or a corresponding derivative of these acids in which the perfluoroalkyl groups contain 1 to 20 C atoms and are straight-chain or branched.
- 20 4. Process according to one or more of Claims 1 to 3,
characterised in that the derivative of bis(fluoroalkyl)phosphinic acid or (fluoroalkyl)phosphonic acid employed is the salt with a mono-, di- or trivalent metal cation.
- 25 5. Process according to Claim 4,
characterised in that the mono-, di- or trivalent metal cation is selected from the group Li^+ , Na^+ , K^+ , Mg^{2+} , Ca^{2+} , Ba^{2+} , Zn^{2+} , Cu^{2+} or Al^{3+} .

6. Process according to one or more of Claims 1 to 3,
characterised in that the derivative of bis(fluoroalkyl)phosphinic acid or (fluoroalkyl)phosphonic acid employed is the salt with a mono- or divalent organic cation.

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7. Process according to Claim 6,
characterised in that the mono- or divalent organic cation is selected from the group tetraalkylammonium, tetraalkylphosphonium, triarylalkylphosphonium, guanidinium, pyrrolidinium, pyridinium, imidazolium, piperazinium or hexamethylenediammonium.

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8. Process according to one of Claims 1 to 3,
characterised in that the derivative of bis(fluoroalkyl)phosphinic acid or (fluoroalkyl)phosphonic acid employed is an ester of bis(fluoroalkyl)phosphinic acid or (fluoroalkyl)phosphonic acid.

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9. Process according to one or more of Claims 1 to 3,
characterised in that the derivative of bis(fluoroalkyl)phosphinic acid or (fluoroalkyl)phosphonic acid employed is the salt with a polycation.

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10. Process according to Claim 9,
characterised in that the polycation is selected from the group of polyammonium cations.

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11. Process according to one or more of Claims 1 to 10,
characterised in that the reaction is carried out in a polar solvent or without a solvent.

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12. Process according to one or more of Claims 1 to 11,
characterised in that the reaction is carried out at a temperature of -20°C to 100°C.

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13. Process according to one or more of Claims 1 to 12,
characterised in that the reaction is carried out with 4- to 100-fold the molar amount of hydrogen fluoride.

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14. Process for the preparation of phosphoranes according to one or more of Claims 1 to 13,
characterised in that the mono- or bis(fluoroalkyl) phosphate formed after the reaction with hydrogen fluoride is reacted with a strong 10 electrophilic reagent or a strong Lewis acid.

15. Process according to Claim 14,
characterised in that the reaction is carried out with an electrophilic reagent or a Lewis acid selected from the group (CH₃)₃SiCl, SO₂Cl₂, SbF₅, AlCl₃, VF₅, SbCl₅, NbF₅, AsF₅, BiF₅, AlF₃ and TaF₅.

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